

REMARKS/ARGUMENTS

Claims 1-8 are canceled.

Claims 9-17 are new.

The claims have been amended to remove multiple dependencies.

The amendments to the claims are supported by the claims, as originally filed, and throughout the specification.

Upon entry of the amendment, Claims 9-17 will be active.

No new matter is believed to have been added. An action on the merits and allowance of the claims is requested.

Respectfully submitted,

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Fractionation of a crude C₄ fraction

5 The present invention relates to a process for fractionating a crude C₄ fraction by extractive distillation using a selective solvent.

10 The term C₄ fraction refers to mixtures of hydrocarbons having predominantly 4 carbon atoms per molecule. C₄ fractions are obtained, for example, in the preparation of ethylene and/or propylene by thermal cracking, for example in steam crackers, in particular naphtha crackers, or FCC plants (fluid catalytic cracking), of a petroleum fraction such as liquefied petroleum gas, naphtha or gas oil. Furthermore, C₄ fractions are obtained in the catalytic dehydrogenation of n-butane and/or n-butene. C₄ fractions generally comprise butane, n-butene, isobutene, 1,3-butadiene and small amounts of other hydrocarbons including 1,2-butadiene, C₅-hydrocarbons and C₄-acetylenes (butynes), in particular 1-butyne (ethylacetylene) and butenyne (vinylacetylene). The 1,3-butadiene content of C₄ fractions from steam crackers is generally from 20 to 70% by weight, in particular from 35 to 65% by weight, while the content of C₄-acetylenes (vinylacetylene and ethylacetylene) generally does not exceed 5% by weight.

20 Owing to the small differences in the relative volatilities of the components, the fractionation of C₄ fractions is a complicated distillation problem. For this reason, fractionation is generally carried out by means of an extractive distillation, i.e. a distillation with addition of a selective solvent (also referred to as extractant) which has a boiling point higher than that of the mixture to be fractionated and which increases the differences in the relative volatilities of the components to be separated.

30 Many processes for fractionating C₄ fractions by means of extractive distillation using selective solvents are known. In all of them, the C₄ fraction in gaseous form is conveyed in countercurrent relative to the liquid selective solvent under suitable thermodynamic conditions, generally at low temperatures, frequently in the range from 20 to 80°C, and moderate pressures, frequently from atmospheric pressure to 6 bar, so that the selective solvent becomes laden with the components from the C₄ fraction for which it has a greater affinity while the components for which the selective solvent has a lower affinity remain in the vapor phase and are taken off as

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overhead stream. The components are subsequently liberated individually or as fractions from the laden solvent stream in one or more further process steps under suitable thermodynamic conditions, i.e. at a temperature which is higher and/or a pressure which is lower than in the first process step.

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In processes for the thermal fractionation of C₄ fractions, C₄-acetylenes present therein present particular problems since they are one of the main causes of apparatus fouling and are prone to spontaneous decomposition within wide concentration ranges.

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For this reason, processes for fractionating C₄ fractions in which the C₄-acetylenes are reacted in a first process step by "front-end hydrogenation" have been developed. Front-end hydrogenation has the further advantage that additional product of value, viz. 1-3-butadiene, is obtained as a result of the hydrogenation of the C₄-acetylenes.

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Such a process is described in Proc.- Ethylene Prod. ~~Conf. 5~~ Conf. 8 (1996), pages 631 to 636. In this process, a high vinylacetylene conversion with a low butadiene loss is achieved using a KLP catalyst, i.e. a catalyst comprising finely divided copper particles on a high-purity γ -aluminum oxide having a defined pore structure as supports, and long catalyst operating lives are also achieved. The upstream selective hydrogenation enables the two-stage butadiene extractive distillation to be simplified to a single-stage process and the apparatus required in the downstream pure distillation to be reduced by one separation column. However, the process has the disadvantage that a separate plant for upstream selective hydrogenation of the acetylene impurities is required.

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US 4,277,313 discloses a further process for recovering 1,3-butadiene in which firstly a selective hydrogenation and subsequently an extractive distillation of the 1,3-butadiene are carried out. The selective hydrogenation can be carried out in the liquid phase or the gas phase, in the presence of catalysts of group VIII of the Periodic Table, for example over a palladium/aluminum oxide catalyst. Extractants mentioned are dimethylformamide or diethylformamide, N-methylpyrrolidone, furfural and acetonitrile. The process has, like the process described above, the disadvantage that the upstream selective hydrogenation requires a separate plant.

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US 6,040,489 discloses a process for separating 1,3-butadiene from a C₄ fraction, in which the C₄ fraction is hydrogenated in a column and selectively extracted by means of a solvent, a stream comprising at least the butanes and butenes is taken off from the column as overhead stream and the solvent laden with butadienes is
5 taken off at the bottom and subsequently separated in a solvent stripping column into a butadiene-containing overhead stream and a solvent-containing bottom stream.

The butadiene-containing overhead stream is separated in a butadiene distillation
10 column into a 1,3-butadiene-containing overhead stream and a 1,2-butadiene-containing bottom stream.

In the process of DE-A 100 22 465.2, a C₄ fraction is subjected to an extractive distillation and a selective hydrogenation over a heterogeneous catalyst in a divid-
15 ing wall column or in thermally coupled columns to give a crude 1,3-butadiene stream.

The known front-end processes for removing C₄-acetylenes from C₄ fractions by selective hydrogenation over heterogeneous catalysts have the disadvantage that
20 not inconsiderable catalyst costs have to be expended and the known catalysts frequently do not have long operating lives. A particularly critical aspect is that when the catalyst becomes exhausted, the entire plant for thermal separation of the C₄ fraction has to be shut down.

25 It is an object of the invention to provide a process for the front-end removal of C₄-acetylenes from C₄ fractions which does not have the abovementioned disadvantages.

We have found that this object is achieved by a process for fractionating a crude C₄
30 fraction comprising butanes, butenes, 1,3-butadiene and small amounts of other hydrocarbons including C₄-acetylenes, 1,2-butadiene and C₅-hydrocarbons by extractive distillation using a selective solvent, wherein the crude C₄ fraction is fed into the middle region of a first extractive distillation column and the selective solvent is fed into the column at a point above that at which the crude C₄ fraction is
35 introduced. A gaseous side stream which comprises the C₄-acetylenes together with 1,3-butadiene, 1,2-butadiene, C₅-hydrocarbons and selective solvent and in

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- which the concentration of the C₄-acetylenes is below the spontaneous decomposition limit is taken off from the first extractive distillation column at a point below the feed point for the crude C₄ fraction and an overhead stream comprising the components of the C₄ fraction which are less soluble than the C₄-acetylenes in the selective solvent is taken off from the top of the first extractive distillation column.

- It has been found that it is economically advantageous and possible in process engineering terms to set operating conditions in an extractive distillation column, in particular in respect of the type of selective solvent, the quantity thereof, temperature, pressure and number of theoretical plates, which enable the C₄-acetylenes, i.e. the components of the C₄ fraction for which the selective solvent has the greatest affinity, to be separated off selectively. This entails process conditions which are unusual for extractive distillation.
- A typical crude C₄ fraction from a naphtha cracker has the following composition in percent by weight:

Propane	0 – 0.5
Propene	0 – 0.5
Propadiene	0 – 0.5
Propyne	0 – 0.5
n-Butane	3 – 10
i-Butane	1 – 3
1-Butene	10 – 20
i-Butene	10 – 30
trans-2-Butene	2 – 8
cis-2-Butene	2 – 6
1,3-Butadiene	35 – 65
1,2-Butadiene	0.1 – 1
Ethylacetylene	0.1 – 2
Vinylacetylene	0.1 – 3
C5	0 – 0.5

- Crude C₄ fractions from naphtha crackers thus comprise predominantly butanes, butenes and 1,3-butadiene. In addition, small amounts of other hydrocarbons are

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present. C₄-acetylenes are frequently present in a proportion of 5% by weight or else up to 2% by weight.

5 Selective solvents suitable for the extractive distillation which has been described at the outset are generally substances or mixtures which have a boiling point higher than that of the mixture to be fractionated and have a greater affinity for conjugated double bonds and triple bonds than for simple double bonds and single bonds, preferably dipolar, particularly preferably dipolar aprotic, solvents. To simplify the choice of apparatus, noncorrosive or relatively noncorrosive substances
10 are preferred.

Examples of suitable selective solvents for the process of the present invention are butyrolactone, nitriles such as acetonitrile, propionitrile, methoxypropionitrile, ketones such as acetone, furfural, N-alkyl-substituted lower aliphatic acid amides,
15 such as dimethylformamide, diethylformamide, dimethylacetamide, diethylacetamide, N-formylmorpholine, N-alkyl-substituted cyclic acid amides (lactams) such as N-alkylpyrrolidones, in particular N-methylpyrrolidone. In general, N-alkyl-substituted lower aliphatic acid amides or N-alkyl-substituted cyclic acid amides are used. Particularly advantageous selective solvents are dimethylformamide,
20 dimide, acetonitrile, furfural and, in particular, N-methylpyrrolidone.

However, it is also possible to use mixtures of these solvents with one another, for example N-methylpyrrolidone with acetonitrile, mixtures of these solvents with cosolvents such as water and/or tert-butyl ethers, for example methyl tert-butyl
25 ether, ethyl tert-butyl ether, propyl tert-butyl ether, n-butyl or isobutyl tert-butyl ether.

A particularly useful solvent is N-methylpyrrolidone, preferably in aqueous solution, in particular with from 8 to 10% by weight of water, particularly preferably
30 8.3% by weight of water.

There are in principle no restrictions regarding the columns which can be used for carrying out the extractive distillation.

35 The C₄ fraction is fed into the column in its middle region and the selective solvent is fed in above the point at which the C₄ fraction is introduced.

The column is provided with separation-active internals. These can be of any known type. Preference is given to one or more trays being located above the feed point for the selective solvent.

- 5 According to the present invention, a side stream comprising the C₄-acetylenes together with 1,3-butadiene, 1,2-butadiene, C₅-hydrocarbons and selective solvent is taken off in gaseous form from the first extractive distillation column, with the column being operated in such a way that the concentration of the C₄-acetylenes in the gaseous side stream is below the spontaneous decomposition limit of these.
- 10 Dilution to below 30 mol% of C₄-acetylenes is generally sufficient for this purpose.

- The selective solvent is present in the gaseous side stream in a proportion corresponding to the thermodynamic equilibrium. The side column is operated as a pure
- 15 rectification column and serves to recover the selective solvent. It has to be operated so that sufficient dilution of the acetylenes to below the spontaneous decomposition region is ensured at every point in the column.

- At the top of the first extractive distillation column, an overhead stream comprising
- 20 the components of the C₄ fraction which are less soluble than the C₄-acetylenes in the selective solvent is taken off. This stream is preferably condensed in a condenser at the top of the column, and part of it is returned as runback to the column while the remainder is worked up further, preferably in a second extractive distillation column. Partial condensation in which the condensed portion serves as run-
- 25 back to the column and the gaseous portion is fed as feed stream to a second extractive distillation column in which separation into raffinate 1 and crude 1,3-butadiene is carried out is particularly advantageous energetically.

- The bottom stream from the first extractive distillation column, which comprises
- 30 predominantly the selective solvent, is preferably used for heat integration into the first extractive distillation column, condensed and recycled to the first extractive distillation column.

- To achieve heat integration, it is possible to cool the hot bottom stream from the
- 35 first extractive distillation column by means of the crude C₄ fraction. In addition or as an alternative, liquid or a substream of the liquid can be taken off from the first

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extractive distillation column at a theoretical plate which is one or more theoretical plates below the point at which the gaseous side stream is taken off, the liquid is heated and/or vaporized by indirect heat exchange with the bottom stream from the first extractive distillation column and is returned to the first extractive distillation column on the same theoretical plate or above this point, with the theoretical plate from which the liquid or liquid substream is taken off being chosen so that the energy requirement for the first extractive distillation column is minimized.

Here, the term raffinate 1 refers in a known manner to a stream comprising butanes and butenes.

The term crude 1,3-butadiene refers to a hydrocarbon mixture in which the desired product 1,3-butadiene is present in a proportion of at least 90% by weight, preferably at least 95% by weight, particularly preferably at least 98% by weight, balance impurities.

With regard to the separation-active internals, what has been said in the context of the first extractive distillation column applies analogously to the second extractive distillation column.

From the second extractive distillation column, a stream is preferably taken off at the top or in the vicinity of the top of the column, condensed in a condenser and the condensate is partly returned as runback to the second extractive distillation column while the remainder is taken off as raffinate 1.

A side stream is taken off from the second extractive distillation column and selective solvent is preferably separated off from this by feeding the side stream into a second short side column where it is separated into selective solvent which is recycled to the second extractive distillation column and an overhead stream which is condensed in a condenser at the top of the column and partly returned as runback to the column while the remainder is taken off as pure 1,3-butadiene.

Heat integration can be achieved in the second extractive distillation column in a manner analogous to the first extractive distillation column by taking off liquid or a substream of the liquid from the second extractive distillation column at a theoretical plate which is one or more theoretical plates below the side offtake, heating

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and/or vaporizing it by indirect heat exchange with the bottom stream from the second extractive distillation column and returning it to the second extractive distillation column on the same theoretical plate or above this, with the theoretical plate from which the liquid or liquid substream is taken off being chosen so that the energy requirement for the second extractive distillation column is minimized.

The invention is illustrated below with the aid of a drawing and an example.

The single figure schematically shows a plant according to the invention for fractionating a C₄ fraction by extractive distillation.

A crude C₄ fraction, stream 1, is fed into the middle region of a first extractive distillation column K I and selective solvent, stream 2, is fed into the column above the point at which the crude C₄ fraction is introduced. A gaseous stream 3 comprising C₄-acetylenes is taken off at a side offtake below the feed point for stream 1 and this is fed to a first side column SK I. In the side column SK I, the stream 3 is separated by distillation into an overhead stream 6 comprising the acetylenes and a bottom stream 7 which comprises selective solvent and is recycled to the first extractive distillation column K I.

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The bottom stream 4 from the first extractive distillation column K I, which comprises predominantly selective solvent, is utilized for heat integration with a liquid stream taken off from the lower region of the first extractive distillation column K I and for preheating the crude C₄ fraction, stream 1, condensed and cooled and returned to the first extractive distillation column K I.

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The overhead stream 5 from the first extractive distillation column is partially condensed in a condenser at the top of the column, the condensate is returned as runback to the column and the gaseous portion is taken off as stream 8.

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In the second extractive distillation column K II, the stream 8 is conveyed in countercurrent to the selective solvent, stream 14, and separated by distillation into an overhead stream 9 which is condensed and part of it is returned as runback to the column K II and the remainder is taken off as raffinate 1, stream 15, and a side stream 10 from which crude 1,3-butadiene is obtained after the solvent has been separated off in a second short side column SK II. In the second side column SK II,

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the side stream 10 is separated into an overhead stream 11 which is condensed in a condenser at the top of the column and part of it is returned as runback to the column and the remainder is taken off as crude 1,3-butadiene, stream 16, and a bottom stream 12 which comprises predominantly the selective solvent and is recycled to the second extractive distillation column K II. At the bottom of the second extractive distillation column K II, a bottom stream 13 is taken off, utilized for heat integration with the liquid stream from the lower region of the second extractive distillation column K II, condensed and subsequently returned as stream 14 to the second extractive distillation column.

Example

A crude C₄ stream, reference 1 in the figure, comprising the following components present in amounts of more than 0.01% by weight in each case:

n-butane	5.75
i-butane	2.45
1-butene	13.89
i-butene	25.65
trans-2-butene	4.44
cis-2-butene	2.96
1,3-butadiene	43.84
1,2-butadiene	0.14
ethylacetylene	0.13 and
vinylacetylene	0.74,

is fed at a flow rate of 32 t/h into an extractive distillation column K I having 28 theoretical plates, numbered from the top, on the 15th theoretical plate.

The column is operated at a pressure at the top of 4.5 bar absolute and a temperature at the top of 58.8°C.

120 t/h of the extractive solvent N-methylpyrrolidone (NMP), containing 8.3% by weight of water, stream 2, is fed in on the upper plate of the extractive distillation column K I.

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A stream having a flow of 224 kg/h, reference 3 in the figure, and comprising the following components present in amounts of more than 0.01% by weight in each case:

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1,3-butadiene	2.45
1,2-butadiene	1.21
ethylacetylene	1.67
vinylacetylene	10.40
water	70.10 and
NMP	14.08,

is taken off from the third theoretical plate.

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The solvent is washed out of this stream in the side column SK I, which is operated as a pure rectification column, by means of the runback. This gives an overhead stream, reference 6 in the figure, comprising the following components present in amounts of greater than 0.01% by weight in each case:

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1,3-butadiene	3.24
1,2-butadiene	1.60
ethylacetylene	2.20
vinylacetylene	13.56
C ₅ -hydrocarbons	0.16 and
water	79.24.

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The extractive solvent N-methylpyrrolidone which has been purchased at considerable cost is completely separated off except for 1 ppm in the overhead stream from the side column and is returned to the first extractive distillation column K I. In this way the acetylenes (ethylacetylene, vinylacetylene) can be removed from the process essentially without losses of the expensive component N-methylpyrrolidone via the overhead stream of the side column SK I..